

The aluminum monostearate is combined with the sesame oil and heated to 125 °C with stirring until a clear yellow solution forms. This mixture is then autoclaved for sterility and allowed to cool. The GH-RH antagonist Peptide 80 is then added aseptically with trituration. Particularly preferred antagonists are salts of low solubility, e.g., pamoate salts and the like. These exhibit long duration of activity.

#### EXAMPLE XI

##### Long Acting Intramuscular (IM) Injectable-Biodegradable Polymer Microcapsules

Microcapsules are made from the following:

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25/75 glycolide/lactide copolymer (0.5 intrinsic viscosity)	99%
[CH <sub>3</sub> (CH <sub>2</sub> ) <sub>6</sub> CO -Tyr <sup>1</sup> , D-Arg <sup>2</sup> , Phe(pCl) <sup>6</sup> , Ala <sup>8</sup> , His <sup>9</sup> , Tyr(Et) <sup>10</sup> , His <sup>11</sup> , Orn <sup>12</sup> , Abu <sup>15</sup> , His <sup>20</sup> , Orn <sup>21</sup> , Nle <sup>27</sup> , D-Arg <sup>28</sup> , Har <sup>29</sup> ]hGH-RH(1-29)NH <sub>2</sub>	(Peptide 96) 1%

15 25 mg of the above microcapsules are suspended in 1.0 mL of the following vehicle:

Dextrose	5.0%
CMC, sodium	0.5%
Benzyl alcohol	0.9%
20 Tween 80	0.1%
Water, purified q.s.	ad 100%

#### EXAMPLE XII

##### Biological Activity in Endocrine and Oncological Assays

25 The peptides of the present invention were tested in assays in vitro and in vivo for their ability to inhibit the hGH-RH(1-29)NH<sub>2</sub> induced GH release. Binding affinities of the compounds to the tumoral GH-RH receptors were also measured. The antitumor activities of the peptides and their inhibitory effects on serum IGF-I and on the tumoral IGF system were evaluated in various cancer models in vivo.

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##### Superfused Rat Pituitary System

The analogs were tested in vitro in a test described earlier (S. Vigh and A.V. Schally, Peptides 5:241-347, 1984) with modification (Z. Rekasi and A.V. Schally, P.N.A.S. 90:2146-2149, 1993).

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Briefly, the cells are preincubated with peptides for 9 minutes (3mL) at various concentrations. Immediately after the incubation, 1 nM hGH-RH(1-29)NH<sub>2</sub> is administered for 3 minutes (1mL) [0 minute response]. To check the duration of the antagonistic effect of the analogue, 1 nM hGH-RH(1-29)NH<sub>2</sub> is applied 30, 60, 90, and 120 minutes later for 3 minutes [30, 60, 90, 120 min responses]. Net integral values of the GH responses are evaluated. GH

40 responses are compared to and expressed as percent of the original GH response induced by 1